

Amendments to the Specification:

Please amend the specification by replacing the paragraph sections under the heading "Related Applications" with the following new paragraph sections:

At page 2, lines 34-37 to page 3, lines 1-8:

R³ is in the 2-, 3- or 4-position and is:

carboxy; (C₁₋₆)alkoxycarbonyl; aminocarbonyl wherein the amino group is optionally substituted by hydroxy, (C₁₋₆)alkyl, hydroxy(C₁₋₆)alkyl, aminocarbonyl(C₁₋₆)alkyl, (C₂₋₆)alkenyl, (C₁₋₆)alkylsulphonyl, trifluoromethylsulphonyl, (C₁₋₆)alkenylsulphonyl, (C₁₋₆)alkoxycarbonyl, (C₁₋₆)alkylcarbonyl, (C₂₋₆)alkenyloxycarbonyl or (C₂₋₆)alkenylcarbonyl and optionally further substituted by (C₁₋₆)alkyl, hydroxy(C₁₋₆)alkyl, aminocarbonyl(C₁₋₆)alkyl or (C₂₋₆)alkenyl; cyano; tetrazolyl; 2-oxo-oxazolidinyl optionally substituted by R¹⁰; 3-hydroxy-3-cyclobutene-1,2-dione-4-yl; 2,4-thiazolidinedione-5-yl; tetrazol-5-ylaminocarbonyl; 1,2,4-triazol-5-yl optionally substituted by R¹⁰; or 5-oxo-1,2,4-oxadiazol-3-yl; or

(C₁₋₄)alkyl **optionally substituted** or ethenyl substituted with any of the substituents listed above for R³ and up to 3 groups R¹² independently selected from:

At page 4, lines 11-19:

A is NR¹¹ or CR⁶R⁷ and B is NR¹¹, O, SO₂ or CR⁸R⁹ and wherein:

each of R⁶, R⁷, R⁸ and R⁹ is independently selected from: hydrogen; (C₁₋₆)alkylthio; halo; trifluoromethyl; azido; (C₁₋₆)alkyl; (C₂₋₆)alkenyl; (C₁₋₆)alkoxycarbonyl; (C₁₋₆)alkylcarbonyl; (C₂₋₆)alkenyloxycarbonyl; (C₂₋₆)alkenylcarbonyl; hydroxy, amino or aminocarbonyl optionally substituted as for corresponding substituents **R¹² as defined** in R³;

(C₁₋₆)alkylsulphonyl; (C₂₋₆)alkenylsulphonyl; or (C₁₋₆)aminosulphonyl wherein the amino group is optionally substituted by (C₁₋₆)alkyl or (C₁₋₆)alkenyl;

or R⁶ and R⁸ together represent a bond and R⁷ and R⁹ are as above defined;

or R⁶ and R⁷ or R⁸ and R⁹ together represent oxo;

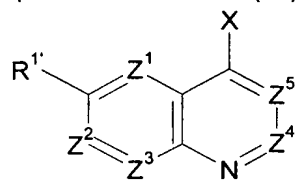
At page 6, lines 3-9:

Preferred examples of R³ include hydrogen; optionally substituted aminocarbonyl; optionally substituted (C~~1-6~~1-4)alkyl; carboxy(C₁₋₄)alkyl; optionally substituted aminocarbonyl(C₁₋₄)alkyl; cyano(C₁₋₄)alkyl; optionally substituted 2-oxo-oxazolidinyl and optionally substituted 2-oxo-oxazolidinyl(C₁₋₄alkyl). More preferred R³ groups are hydrogen; CONH₂; 1-hydroxyalkyl e.g. CH₂OH, CH(OH)CH₂CN; CH₂CO₂H; CH₂CONH₂; 1,2-dihydroxyalkyl e.g. CH(OH)CH₂OH; CH₂CN; 2-oxo-oxazolidin-5-yl and 2-oxo-oxazolidin-5-yl(C₁₋₄alkyl).

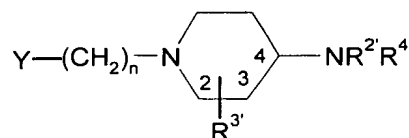
At page 8, lines 32-36 to page 9, lines 1-31:

In a further aspect of the invention there is provided a process for preparing compounds of formula (I), and pharmaceutically acceptable derivatives thereof, which process comprises:

reacting a compound of formula (IV) with a compound of formula (V):



(IV)



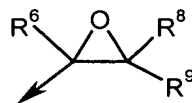
(V)

wherein Z^1 , Z^2 , Z^3 , Z^4 , Z^5 and n are as defined in formula (I); $R^{1'}$, $R^{2'}$, $R^{3'}$ and $R^{4'}$ are R^1 , R^2 , R^3 and R^4 as defined in formula (I) or groups convertible thereto;

and X and Y may be the following combinations:

- (i) X is $A'-COW$, Y is H and n is 0;
- (ii) X is $CR^6=CR^8R^9$, Y is H and n is 0;
- (iii) X is oxirane, Y is H and n is 0;
- (iv) X is $N=C=O$ and Y is H ;
- (v) X is NH_2 and Y is CO_2W ;
- (vi) one of X and Y is CO_2R^Y and the other is $CH_2CO_2R^X$;
- (vii) X is CHR^6R^7 and Y is CR^8O ;
- (viii) X is $CR^6=PR^Z_3$ and Y is CR^8O ;
- (ix) X is CR^6O and Y is $CR^8=PR^Z_3$;
- (x) one of X and Y is COW and the other is $NHR^{11'}$ or NCO ;
- (xi) X is CR^6O and Y is $NHR^{11'}$ or X is $NHR^{11'}$ and Y is CR^8O ;
- (xii) X is $NHR^{11'}$ and Y is CR^8R^9W ;
- (xiii) X is CR^6R^7W and Y is $NR^{11'}$ or O ; or
- (xiv) X is $CR^6R^7SO_2W$ and Y is H and $n=0$;
- (xv) X is $NR^{11'}$ and Y is SO_2W ;

in which W is a leaving group, e.g. halogen; R^X and R^Y are (C_{1-6}) alkyl; R^Z is aryl or (C_{1-6}) alkyl; A' and $NR^{11'}$ are A and NR^{11} as defined in formula (I), or groups convertible thereto; and oxirane is:



wherein R^6 , R^8 and R^9 are as defined in formula (I);

and thereafter optionally or as necessary converting A' , $R^{1'}$, $R^{2'}$, $R^{3'}$, $R^{4'}$ and $NR^{11'}$ to A , R^1 , R^2 , R^3 , R^4 and NR^{11} ; converting $A-B$ to other $A-B$, interconverting R^1 , R^2 , R^3 and/or R^4 , and/or forming a pharmaceutically acceptable derivative thereof.